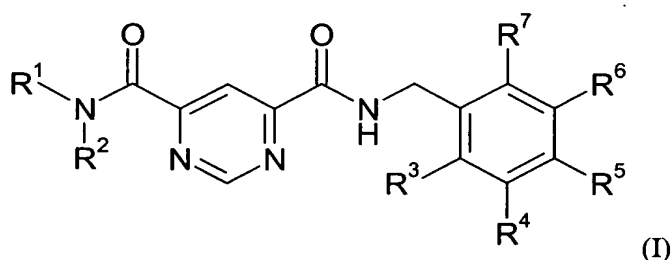


What is claimed is:

1. A compound of formula I



wherein

R<sup>1</sup> is

hydrogen atom or  $-(C_1-C_6)$ -alkyl,

R<sup>2</sup> is

$-(C_1-C_6)$ -alkyl that is substituted, once, twice or three times, by

$-C(O)-O-R^8$ ,

$-(C_1-C_6)$ -alkyl- $O-R^8$ ,

$-(C_6-C_{14})$ -aryl that is substituted, once, twice or three times, independently of each other, by R<sup>11</sup> or

Het that is a saturated or unsaturated monocyclic or bicyclic, 3- to 10-membered heterocyclic ring system which contains 1, 2 or 3

identical or different ring heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur and is unsubstituted or substituted, once or more than once, by R<sup>13</sup>,

R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are identical or different and are, independently of each other,

hydrogen

halogen,

$-(C_1-C_6)$ -alkyl, in which alkyl is unsubstituted or substituted, once, twice or three times, by halogen,

$-O-(C_1-C_6)$ -alkyl, in which alkyl is unsubstituted or substituted, once, twice or three times, by halogen, or

$-S-(C_1-C_6)$ -alkyl,

R<sup>8</sup> is

hydrogen atom, or  
 -(C<sub>1</sub>-C<sub>6</sub>)-alkyl,

5

R<sup>11</sup> is

-(C<sub>2</sub>-C<sub>6</sub>)-alkyl-C(O)-O-R<sup>8</sup>,  
 -O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl-C(O)-O-R<sup>8</sup>,  
 -NR<sup>14</sup>R<sup>15</sup>,  
 10 -(CH<sub>2</sub>)<sub>k</sub>-NR<sup>9</sup>R<sup>10</sup>,  
 -O-(C<sub>2</sub>-C<sub>6</sub>)-alkyl-NR<sup>9</sup>R<sup>10</sup>, or  
 -NR<sup>8</sup>-C(O)-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, in which alkyl is unsubstituted or substituted, once,  
 twice or three times, by R<sup>12</sup>,

15 R<sup>9</sup> and R<sup>10</sup> are identical or different and are, independently of each other,

hydrogen atom, or  
 -(C<sub>1</sub>-C<sub>6</sub>)-alkyl, or  
 taken together with the nitrogen atom to which they are attached form a 5-, 6- or  
 7-membered saturated azaheterocyclyl ring wherein one or two further carbon  
 20 atoms thereof are optionally replaced by a heteroatom that is an oxygen, sulfur or  
 nitrogen atom, and wherein the nitrogen atom is optionally unsubstituted or  
 substituted by (C<sub>1</sub>-C<sub>6</sub>)-alkyl,

k is

25 2, 3, 4 or 5,

R<sup>12</sup> is

halogen,  
 cyano,  
 30 nitro,  
 hydroxyl,  
 amino,  
 -C(O)-O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, or  
 -C(O)-OH,

35

R<sup>13</sup> is

- halogen,
- cyano,
- nitro,
- 5 hydroxyl,
- amino,
- C(O)-O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl,
- C(O)-OH,
- (C<sub>1</sub>-C<sub>6</sub>)-alkyl that is unsubstituted or substituted, once, twice or three times, by
- 10 halogen,
- O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, where alkyl is unsubstituted or substituted, once, twice or three
- times, by halogen,
- pyridyl, or
- phenyl that is unsubstituted or substituted, once or more than once and
- 15 independently of each other, by a radical from the series halogen, (C<sub>1</sub>-C<sub>6</sub>)-
- alkoxy and (C<sub>1</sub>-C<sub>6</sub>)-alkyl, and

R<sup>14</sup> and R<sup>15</sup> together with the nitrogen atom to which they are attached form

- a 5-, 6- or 7-membered saturated azaheterocyclyl ring wherein one or two further
- 20 carbon atoms thereof are optionally replaced by a heteroatom that is oxygen,
- sulfur or nitrogen, and wherein each nitrogen atom thereof is optionally
- independently unsubstituted or substituted by (C<sub>1</sub>-C<sub>6</sub>)-alkyl, or

- stereoisomer thereof, a mixture of stereoisomers thereof in any ratio, or physiologically
- 25 tolerable salt thereof.

2. The compound according to claim 1, wherein

R<sup>2</sup> is

- (C<sub>1</sub>-C<sub>4</sub>)-alkyl, where alkyl is substituted, once, twice or three times, by
- 30 -C(O)-O-R<sup>8</sup>,
- (C<sub>1</sub>-C<sub>4</sub>)-alkyl-O-R<sup>8</sup>,
- phenyl that is substituted, once, twice or three times, independently of
- each other, by R<sup>11</sup>, or
- Het that is azepine, azetidine, aziridine, benzimidazole, benzo[1,4]dioxin,
- 35 1,3-benzodioxole, benzofuran, 4H-benzo[1,4]oxazine,

benzoxazole, benzothiazole, benzothiophene, quinazoline,  
 quinoline, quinoxaline, chroman, cinnoline, oxirane,  
 1,2-diazepine, 1,3-diazepine, 1,4-diazepine, 1,4-dioxin, dioxole,  
 furan, imidazole, indazole, indole, isoquinoline, isochroman,  
 5 isoindole, isoxazole, isothiazole, 1,2-oxazine, 1,3-oxazine,  
 1,4-oxazine, oxazole, phthalazine, piperidine, pyran, pyrazine,  
 pyrazole, pyridazine, pyridine, pyrimidine, pyridoimidazole,  
 pyridopyridine, pyridopyrimidine, pyrrol, tetrazole, 1,2-thiazine,  
 1,3-thiazine, 1,4-thiazine, thiazole, thiophene, thiopyran,  
 10 1,2,3-triazine, 1,2,4-triazine, 1,3,5-triazine, 1,2,3-triazole or 1,2,4-  
 triazole, and Het is unsubstituted or substituted, once, twice or  
 three times, independently of each other, by R<sup>13</sup>

R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are identical or different and are  
 15 hydrogen atom,  
 halogen,  
 -(C<sub>1</sub>-C<sub>6</sub>)-alkyl, in which alkyl is unsubstituted or substituted, once, twice or three  
 times, by halogen, or  
 -O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, in which alkyl is unsubstituted or substituted, once, twice or  
 20 three times, by halogen,

R<sup>8</sup> is  
 hydrogen atom, or  
 -(C<sub>1</sub>-C<sub>4</sub>)-alkyl,

25 R<sup>11</sup> is  
 -(C<sub>2</sub>-C<sub>4</sub>)-alkyl-C(O)-O-R<sup>8</sup>,  
 -O-(C<sub>1</sub>-C<sub>4</sub>)-alkyl-C(O)-O-R<sup>8</sup>,  
 -N R<sup>14</sup>R<sup>15</sup>, wherein R<sup>14</sup> and R<sup>15</sup> taken together with the nitrogen atom to which  
 30 they are attached form imidazolidine, isothiazolidine, isoxazolidine,  
 morpholine, piperazine, piperidine, pyrazine, pyrazolidine, pyrrolidine,  
 tetrazine or thiomorpholine, and wherein each nitrogen atom thereof is  
 optionally independently unsubstituted or substituted by (C<sub>1</sub>-C<sub>4</sub>)-alkyl,  
 -(CH<sub>2</sub>)<sub>k</sub>-N R<sup>9</sup>R<sup>10</sup>,  
 35 -O-(C<sub>2</sub>-C<sub>4</sub>)-alkyl-NR<sup>9</sup>R<sup>10</sup>, or

-NH-C(O)-(C<sub>1</sub>-C<sub>4</sub>)-alkyl, wherein the alkyl is unsubstituted or substituted, once, twice or three times, by R<sup>12</sup>,

R<sup>9</sup> and R<sup>10</sup> are identical or different and are, independently of each other,

5        hydrogen atom, or  
      -(C<sub>1</sub>-C<sub>4</sub>)-alkyl, or  
      taken together with the nitrogen atom to which they are attached form  
      imidazolidine, isothiazolidine, isoxazolidine, morpholine, piperazine, piperidine,  
      pyrazine, pyrazolidine, pyrrolidine, tetrazine or thiomorpholine, and wherein the  
10        nitrogen atom is optionally unsubstituted or substituted by -(C<sub>1</sub>-C<sub>4</sub>)-alkyl,

k is

2, 3 or 4, and

15        R<sup>13</sup> is

      halogen,  
      amino,  
      -C(O)-O-(C<sub>1</sub>-C<sub>4</sub>)-alkyl,  
      -C(O)-OH,  
20        -(C<sub>1</sub>-C<sub>6</sub>)-alkyl that is unsubstituted or substituted, once, twice or three times, by  
          halogen,  
      -O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, wherein the alkyl is unsubstituted or substituted, once, twice or  
          three times, by halogen,  
      pyridyl, or  
25        phenyl that is unsubstituted or substituted, once or more than once and  
          independently of each other, by a radical from the series halogen,  
          -(C<sub>1</sub>-C<sub>4</sub>)-alkoxy and -(C<sub>1</sub>-C<sub>4</sub>)-alkyl.

3.        The compound according to claim 1, wherein

30        R<sup>1</sup> is

hydrogen,

R<sup>2</sup> is

-(C<sub>1</sub>-C<sub>2</sub>)-alkyl that is substituted, once, twice or three times, by

phenyl that is substituted, once, twice or three times, independently of each other, by  $R^{11}$ , or

Het that is furan, imidazole, isothiazole, isoxazole, oxazole, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, thiazole, thiophene, 1,2,3-triazole or 1,2,4-triazole, and Het is unsubstituted or substituted, once, twice or three times, independently of each other, by  $R^{13}$ ,

$R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^7$  are identical or different and are, independently of each other,

hydrogen,  
halogen,  
methyl,  
trifluoromethyl,  
methoxy, or  
trifluoromethoxy,

$R^8$  is

hydrogen atom, or  
 $-(C_1-C_4)$ -alkyl,

$R^{11}$  is

$-(C_2-C_4)$ -alkyl-C(O)-O- $R^8$ ,  
 $-O-(C_1-C_4)$ -alkyl-C(O)-O- $R^8$ ,  
 $-N R^{14} R^{15}$ , wherein  $R^{14}$  and  $R^{15}$  taken together with the nitrogen atom to which they are attached form imidazolidine, isothiazolidine, isoxazolidine, morpholine, piperazine, piperidine, pyrazine, pyrazolidine, pyrrolidine, tetrazine or thiomorpholine, and wherein each nitrogen atom thereof is optionally independently unsubstituted or substituted by  $(C_1-C_4)$ -alkyl,  
 $-(CH_2)_k-N R^9 R^{10}$ ,  
 $-O-(C_2-C_4)$ -alkyl-N $R^9 R^{10}$ , or  
 $-NH-C(O)-(C_1-C_4)$ -alkyl, wherein the alkyl is unsubstituted or substituted, once, twice or three times, by  $R^{12}$ ,

$R^9$  and  $R^{10}$  are identical or different and are, independently of each other,

hydrogen atom, or

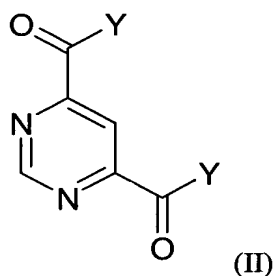
- 5         $-(C_1-C_4)$ -alkyl, or  
          taken together with the nitrogen atom to which they are attached form  
          imidazolidine, isothiazolidine, isoxazolidine, morpholine, piperazine, piperidine,  
          pyrazine, pyrazolidine, pyrrolidine, tetrazine or thiomorpholine, and wherein the  
          nitrogen atom is optionally unsubstituted or substituted by  $-(C_1-C_4)$ -alkyl,
- k is
- 2, 3 or 4,
- 10        $R^{12}$  is
- halogen,  
           $-C(O)-O-(C_1-C_4)$ -alkyl, or  
           $-C(O)-OH$ , and
- 15        $R^{13}$  is
- halogen,  
          amino,  
           $-C(O)-O-(C_1-C_4)$ -alkyl,  
           $-C(O)-OH$ ,
- 20        $-(C_1-C_4)$ -alkyl that is unsubstituted or substituted, once, twice or three times, by  
          halogen,  
           $-O-(C_1-C_4)$ -alkyl, wherein the alkyl is unsubstituted or substituted, once, twice or  
          three times, by halogen,  
          pyridyl, or
- 25       phenyl that is unsubstituted or substituted, once or more than once and  
          independently of each other, by a radical from the series halogen,  
           $-(C_1-C_4)$ -alkoxy and  $-(C_1-C_4)$ -alkyl.
- 30       4.       A method for the prophylaxis or therapy of a patient having or subject to a disease  
          whose course involves a detrimental increase in the activity of matrix  
          metalloproteinase 13, comprising administering to said patient a therapeutically  
          effective amount of a compound according to claim 1.
- 35       5.       A method for the prophylaxis or therapy of a patient having or subject to a disease  
          whose course involves a detrimental increase in the activity of matrix

metalloproteinase 13, comprising administering to said patient a therapeutically effective amount of a compound according to claim 2.

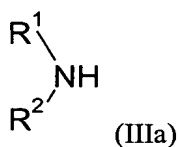
- 5 6. A method for the prophylaxis or therapy of a patient having or subject to a disease whose course involves a detrimental increase in the activity of matrix metalloproteinase 13, comprising administering to said patient a therapeutically effective amount of a compound according to claim 3.

- 10 7. A process for preparing the compound of formula I according to claim 1, comprising

a) reacting a compound of formula II

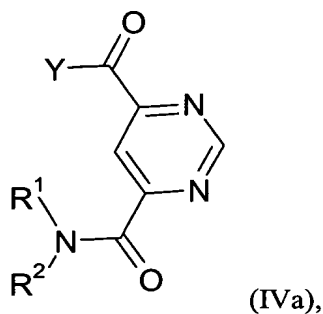


- 15 wherein Y is  
halogen, hydroxyl or C<sub>1</sub>-C<sub>4</sub>-alkoxy, or forms, together with the carbonyl group, an active ester or a mixed anhydride,  
with a compound of formula IIIa

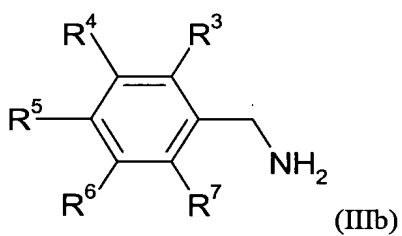


20 wherein R<sup>1</sup> and R<sup>2</sup>, have the meanings given in the compound of formula I, to form a compound of formula IVa



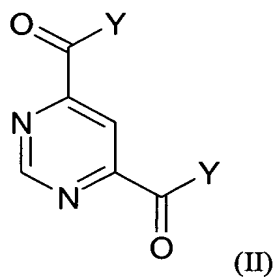


- b) reacting the compound of formula IVa with a compound of formula IIIb



wherein  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^7$  have the meanings given in the compound of formula I, to form the compound of formula I.

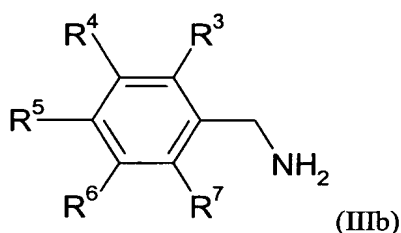
8. A process for preparing the compound of formula I according to claim 1, comprising
- a) reacting a compound of formula II



wherein Y is

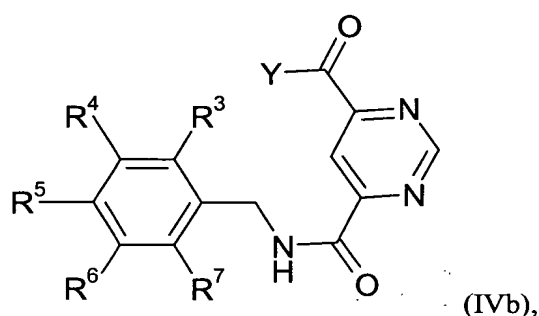
halogen, hydroxyl or  $C_1$ - $C_4$ -alkoxy, or forms, together with the carbonyl group, an active ester or a mixed anhydride,

with a compound of formula IIIb



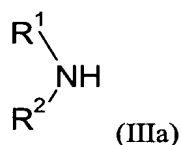
wherein  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^7$  have the meanings given in the compound of formula I,

5 to form a compound of formula IVb



b) reacting the compound of formula IVb with a compound of formula IIIa

10



wherein  $R^1$  and  $R^2$ , have the meanings given in the compound of formula I,  
to form the compound of formula I.

15

9. A pharmaceutical preparation comprising a pharmaceutically effective amount of at least one compound of formula I according to claim 1 and a pharmaceutically suitable and physiologically tolerated carrier.

20 10. A use of the compound according to claim 1 for the prophylaxis or therapy of a patient having or subject to a disease that involves a detrimental increase in the activity of matrix metalloproteinase 13, comprising administering to the patient a pharmaceutically effective amount of at least one compound of formula I.

11. The use according to claim 10 wherein the disease is a degenerative joint disease, or disease of the connective tissue, chronic disease of the locomotory apparatus or cancer disease such as breast cancer.